

U. S. NUCLEAR REGULATORY COMMISSION

Docket No. NRC-2008-0368

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: U. S. Nuclear Regulatory Commission (NRC).

ACTION: Notice of pending NRC action to submit an information collection request to the Office of Management and Budget (OMB) and solicitation of public comment.

SUMMARY: The NRC is preparing a submittal to OMB for review of continued approval of information collections under the provisions of the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35).

Information pertaining to the requirement to be submitted:

1. *The title of the information collection:* Registration Certificate In-Vitro Testing with ByProduct Material under General License.
2. *Current OMB approval number:* 3150-0038.
3. *How often the collection is required:* There is a one-time submittal of information to receive a validated copy of NRC Form 483 with an assigned registration number. In addition, any changes in the information

reported on NRC Form 483 must be reported in writing to the Commission within 30 days after the effective date of such change.

4. *Who is required or asked to report:* Any physician, veterinarian in the practice of veterinary medicine, clinical laboratory or hospital which desires a general license to receive, acquire, possess, transfer, or use specified units of byproduct material in certain in vitro clinical or laboratory tests.
5. *The number of annual respondents:* 85.
6. *The number of hours needed annually to complete the requirement or request:* 12.4 hours (Record keeping: 1.13 hours + Reporting: 2 hours NRC licensees and 9.3 hours Agreement State licensees).
7. *Abstract:* Section 31.11 of 10 CFR establishes a general license authorizing any physician, clinical laboratory, veterinarian in the practice of veterinary medicine, or hospital to possess certain small quantities of byproduct material for in vitro clinical or laboratory tests not involving the internal or external administration of the byproduct material or the radiation there from to human beings or animals. Possession of byproduct material under 10 CFR 31.11 is not authorized until the physician, clinical laboratory, veterinarian in the practice of veterinary medicine, or hospital has filed NRC Form 483 and received from the

Commission a validated copy of NRC Form 483 with a registration number.

Submit, by (**insert date** 60 days after publication in the Federal Register), comments that address the following questions:

1. Is the proposed collection of information necessary for the NRC to properly perform its functions? Does the information have practical utility?
2. Is the burden estimate accurate?
3. Is there a way to enhance the quality, utility, and clarity of the information to be collected?
4. How can the burden of the information collection be minimized, including the use of automated collection techniques or other forms of information technology?

